

Serpine2 Cas9-KO Strategy

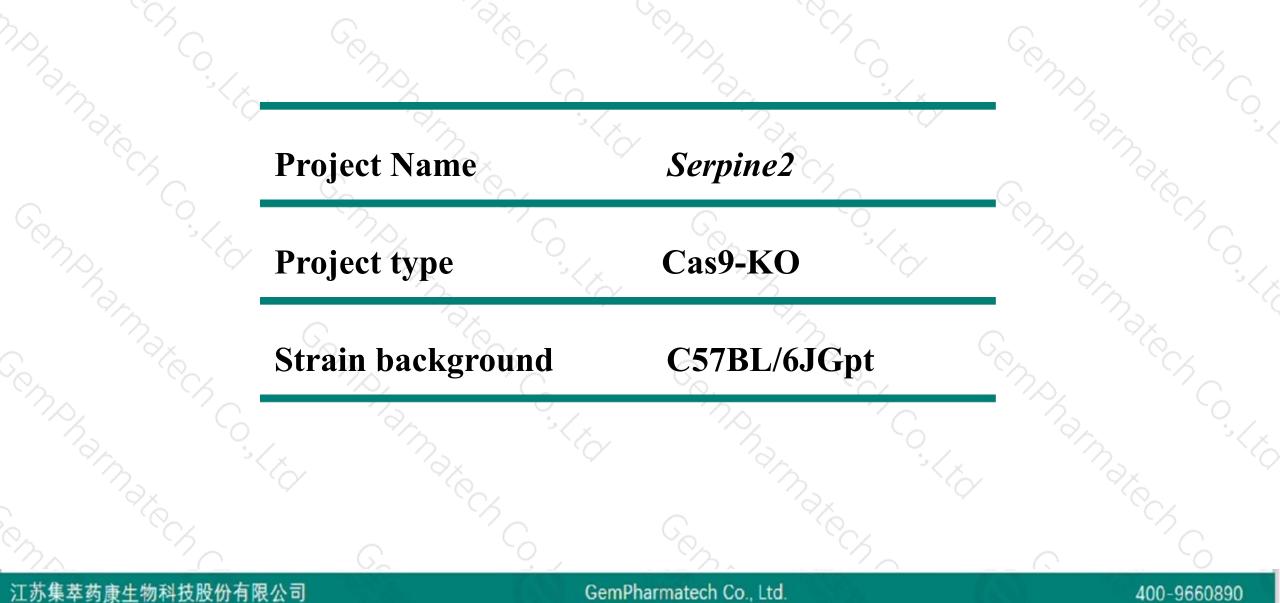
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Reviewer: Shanhong Tao

Design Date: 2021-3-31

Project Overview

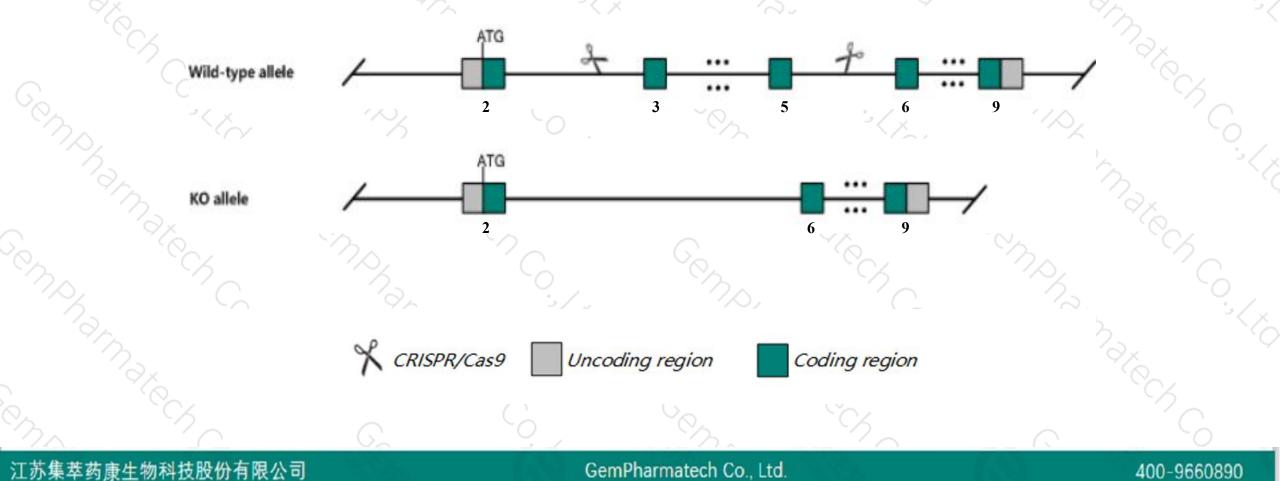




Knockout strategy



This model will use CRISPR/Cas9 technology to edit the Serpine2 gene. The schematic diagram is as follows:





> The Serpine2 gene has 6 transcripts. According to the structure of Serpine2 gene, exon3-exon5 of Serpine2-201(ENSMUST00000027467.10) transcript is recommended as the knockout region. The region contains 625bp coding sequence. Knock out the region will result in disruption of protein function.

➤ In this project we use CRISPR/Cas9 technology to modify *Serpine2* gene. The brief process is as follows: CRISPR/Cas9 system were microinjected into the fertilized eggs of C57BL/6JGpt mice.Fertilized eggs were transplanted to obtain positive F0 mice which were confirmed by PCR and sequencing. A stable F1 generation mouse model was obtained by mating positive F0 generation mice with C57BL/6JGpt mice.



- > According to the existing MGI data, mice homozygous for a targeted mutation of this gene are viable and healthy but develop epileptic activity as well as reduced theta burst-induced LTP and NMDA receptor-mediated synaptic transmission in the CA1 field of the hippocampus; notably, homozygous mutant males are infertile.
- > The *Serpine2* gene is located on the Chr1. If the knockout mice are crossed with other mice strains to obtain double gene positive homozygous mouse offspring, please avoid the two genes on the same chromosome.
- ➤ This strategy is designed based on genetic information in existing databases.Due to the complexity of biological processes,all risk of the gene knockout on gene transcription, RNA splicing and protein translation cannot be predicted at the existing technology level.

Gene information (NCBI)



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400-9660890

Serpine2 serine (or cysteine) peptidase inhibitor, clade E, member 2 [Mus musculus (house mouse)]

Gene ID: 20720, updated on 13-Mar-2020

Summary

Official Symbol	Serpine2 provided by MGI
Official Full Name	serine (or cysteine) peptidase inhibitor, clade E, member 2 provided byMGI
Primary source	MGI:MGI:101780
See related	Ensembl:ENSMUSG00000026249
Gene type	protein coding
RefSeq status	VALIDATED
Organism	Mus musculus
Lineage	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Euarchontoglires; Glires; Rodentia; Myomorpha;
	Muroidea; Muridae; Murinae; Mus; Mus
Also known as	B230326M24Rik, PAI-1, PI-7, PI7, PN-1, Spi4
Expression	Broad expression in ovary adult (RPKM 75.8), frontal lobe adult (RPKM 44.2) and 16 other tissues See more
Orthologs	human all

Transcript information (Ensembl)



The gene has 6 transcripts, all transcripts are shown below:

Name	Transcript ID	bp	Protein	Biotype	CCDS	UniProt	Flags
Serpine2-201	ENSMUST00000027467.10	2210	<u>397aa</u>	Protein coding	CCDS15092	Q07235 Q543R5	TSL:1 GENCODE basic APPRIS P1
Serpine2-204	ENSMUST00000190724.1	822	<u>232aa</u>	Protein coding		A0A087WQM1	CDS 3' incomplete TSL:5
Serpine2-203	ENSMUST00000189793.6	696	<u>232aa</u>	Protein coding	84	A0A087WQ70	CDS 5' and 3' incomplete TSL:5
Serpine2-206	ENSMUST00000191529.1	3393	No protein	Retained intron	10 11	10210	TSL:NA
Serpine2-202	ENSMUST00000153862.1	2315	No protein	Retained intron	17	171	TSL:1
Serpine2-205	ENSMUST00000191026.1	1129	No protein	Retained intron	-	-	TSL:NA

The strategy is based on the design of Serpine2-201 transcript, the transcription is shown below:

< Serpine2-201 protein coding

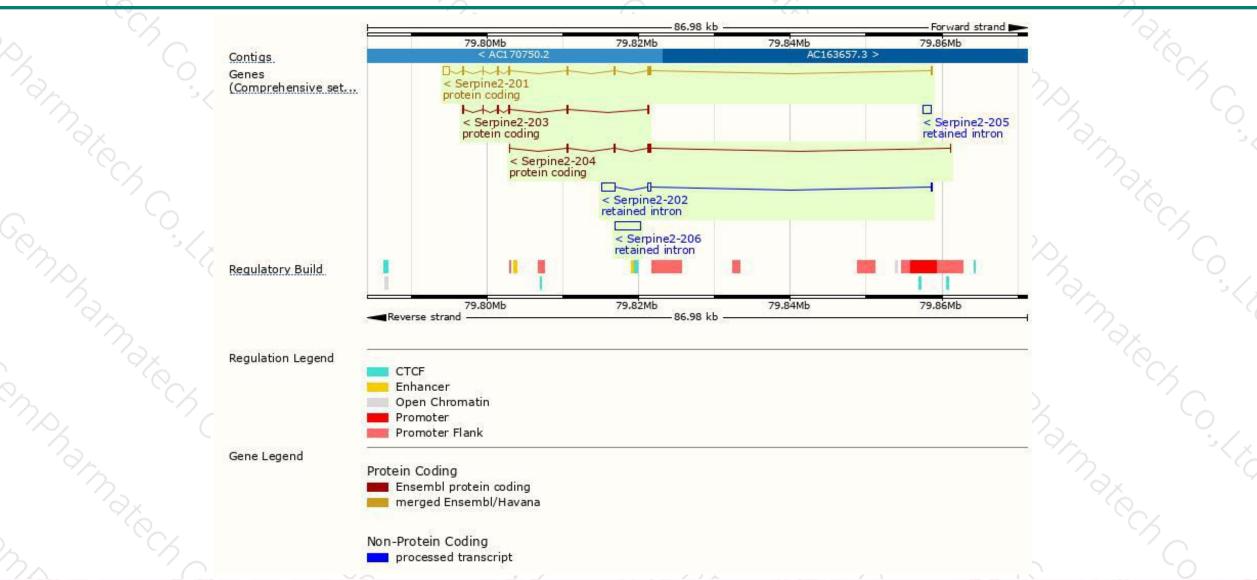
Reverse strand

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64.50 kb

Genomic location distribution



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Protein domain



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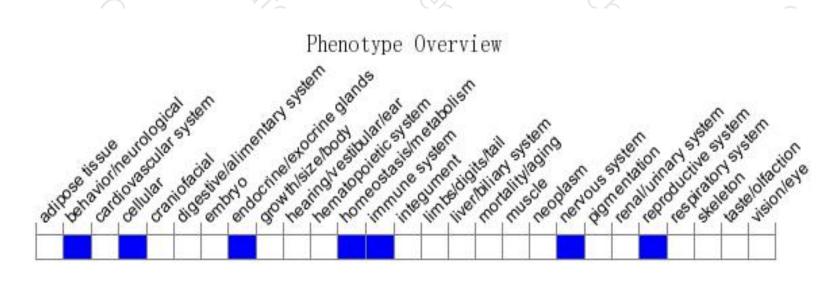
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Mouse phenotype description(MGI)





Phenotypes affected by the gene are marked in blue. Data quoted from MGI database(http://www.informatics.jax.org/).

According to the existing MGI data, mice homozygous for a targeted mutation of this gene are viable and healthy but develop epileptic activity as well as reduced theta burst-induced LTP and NMDA receptor-mediated synaptic transmission in the CA1 field of the hippocampus; notably, homozygous mutant males are infertile.

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If you have any questions, you are welcome to inquire. Tel: 400-9660890



